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Preliminary Testing of a Prototype Portable X-Ray Fluorescence Spectrometer

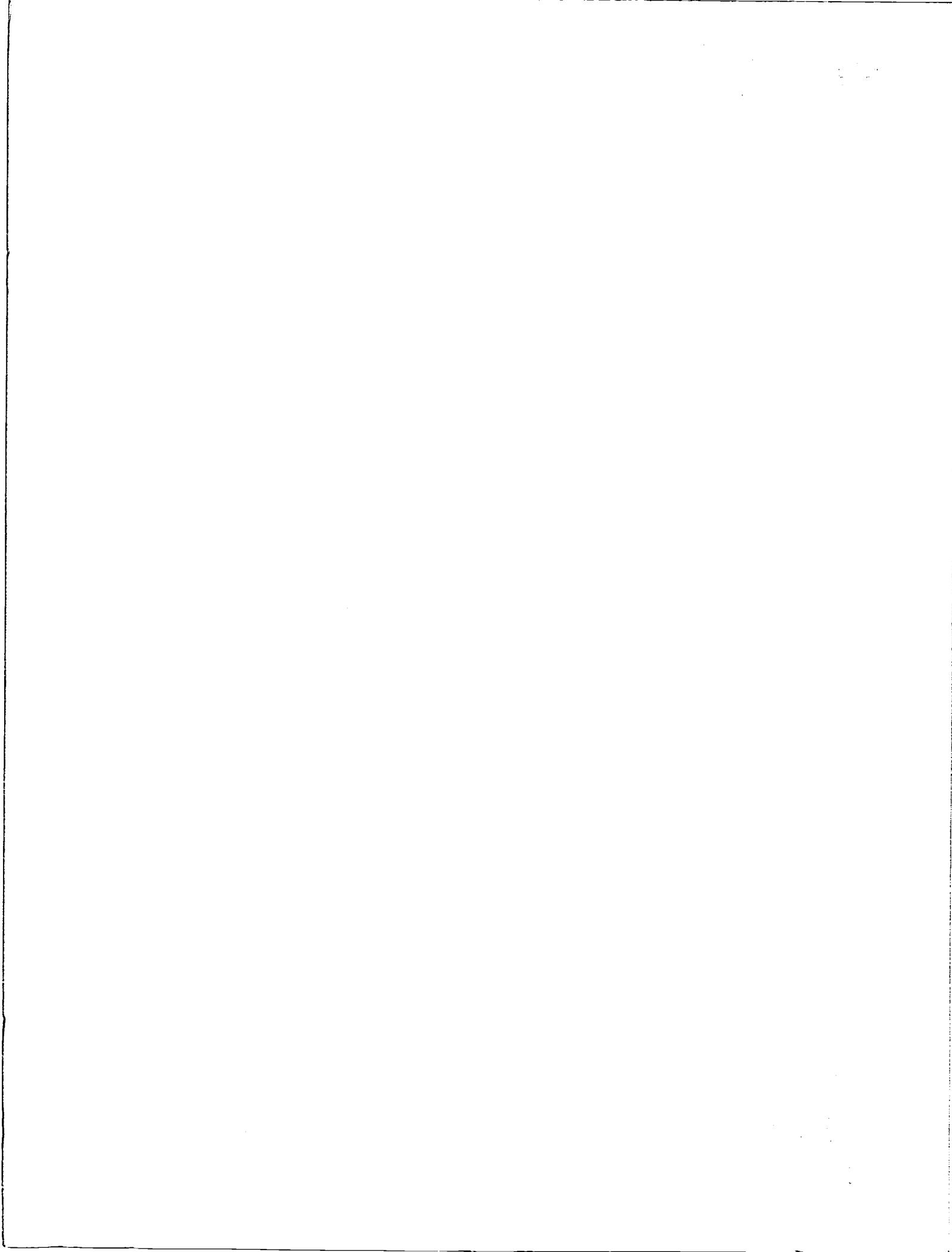
**By Lowell L. Patten, Neal B. Anderson,
and John J. Stevenson**

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UNITED STATES DEPARTMENT OF THE INTERIOR



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UNITED STATES DEPARTMENT OF THE INTERIOR

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PRELIMINARY TESTING OF A PROTOTYPE PORTABLE X-RAY FLUORESCENCE SPECTROMETER

By Lowell L. Patten,¹ Neal B. Anderson,² and John J. Stevenson²

ABSTRACT

The Federal Bureau of Mines participated with the National Aeronautics and Space Administration and Martin Marietta Aerospace in developing, building, and testing a portable X-ray fluorescence spectrometer for use as an analyzer in mineral-resource investigative work. The prototype battery-powered spectrometer, measuring 11 by 12 by 5 inches and weighing only about 15 pounds, was designed specifically for field use. The spectrometer has two gas-proportional counters and two radioactive sources, ¹⁰⁹Cd and ⁵⁵Fe. Preliminary field and laboratory tests on rock specimens and rock pulps have demonstrated the capability of the spectrometer to detect 33 elements, to date. Characteristics of the system present some limitations, however, and further improvements are recommended.

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INTRODUCTION

The principles of X-ray fluorescence spectrometry long have been known and studied intensively by numerous investigators (see bibliography). Laboratory model spectrometers have been available commercially for many years. The feasibility of a portable X-ray fluorescence spectrometer (PXRFS) was investigated in the 1960's, and several companies developed portable analyzers, some using X-ray generators and others using radioactive isotope sources. In general, however, these analyzers were heavy, cumbersome, complex to use, and lacking in versatility. Complications included the need to cool the X-ray generators, the use of filters to increase discrimination of the detectors, and the fact that most of these analyzers would detect only one or no more than a few elements at one time.

An improved portable spectrometer would greatly facilitate the Federal Bureau of Mines mineral-land assessment work, would substantially assist mineral exploration work in general, and could benefit both Government and industry as an aid in mineral identification and analysis.

This report summarizes results of a twofold project to apply principles of X-ray fluorescence spectrometry to mineral identification and element quantification. The first part of the project was to construct a portable, energy dispersive, X-ray fluorescence spectrometer. The second part was to test the prototype instrument in laboratory and field situations to determine its operating characteristics, its response, and its usefulness in mineral-resource investigative work.

Instrument response was observed and recorded in the Planetary Geology Laboratory of Martin Marietta Aerospace near Denver, Colo., in the Bureau's Intermountain Field Operations Center at the Denver (Colo.) Federal Center, in the mountains west of Denver, and near Tucson, Ariz.

During testing, a limited effort was made to compare the PXRFS capabilities with those of emission spectrometers. The PXRFS can be used in conjunction with, but not as a substitute for, a laboratory emission spectrometer.

History

The idea of a spectrometer that could be carried easily in the field and could quickly identify and quantify many elements with little or no sample preparation only recently became technologically possible.

In the 1970's, Martin Marietta (MM), under contract with National Aeronautic and Space Administration (NASA), developed a miniature X-ray fluorescence spectrometer, which employed ^{55}Fe and ^{109}Cd isotopes as X-ray sources, for use on the Viking Project Mars Lander.

The senior author of this report suggested that the Mars Lander technology might be followed in developing a portable spectrometer that could be useful as a field instrument in mineral exploration. Subsequently, Bureau personnel consulted with representatives of MM and NASA, and in 1975 a contract was negotiated for developing and constructing a portable X-ray fluorescence spectrometer. The work was funded mostly by the Bureau and it also provided a list of elements of interest in mineral exploration. Suggestions for design to facilitate field and laboratory use were contributed by both the Bureau and NASA.

The PXRFS was completed in mid-1980; subsequently, several minor adjustments and microprocessor program modifications were made. A testing program began in late 1980 and continued intermittently until the spring of 1981. A Nuclear Regulatory Commission (NRC) licensing (see appendix C for license requirements) problem delayed transfer of the spectrometer custody from the contractor to the

Bureau until July 1981 when the Bureau's source license was modified to include the PXRFS and its radioactive isotopes.

Previous Bureau of Mines Work

The Bureau of Mines has done considerable work in laboratory and field

applications of X-ray fluorescence as shown by citations in the bibliography. Most of this research advanced the general technology but was directed mainly toward special applications, and, in particular, toward high quantitative accuracy in the laboratory.

ACKNOWLEDGMENTS

Those who technically contributed the most to the current project were Benton C. Clark, senior research scientist, and Ludwig Wolfert, staff engineer, both of Martin Marietta. Also instrumental in the development of the PXRFS

were Warren C. Kelliher, contract manager, and Charles R. Eastwood, manager, environmental projects, both from NASA, and Sheldon P. Wimpfen, chief mining engineer, and Lee R. Rice, geologist, of the Bureau of Mines.

X-RAY THEORY AND ITS APPLICATION TO THE PORTABLE SPECTROMETER

X-rays can be generated by electrical apparatus or originate as gamma rays during the decay of radioactive isotopes of elements. The origin of characteristic X-ray fluorescence can briefly be described as follows: When sufficient energy is introduced into the atom, by X-rays or gamma rays, an electron is displaced from one of the inner shells. The atom is then in an excited (ionized) state. The place of the missing electron is filled immediately by an electron from a neighboring outer shell whose place, in turn, is filled by an electron from the next outer shell. The electron from a high energy level (outer shell) enters a lower energy level (inner shell), emitting excess energy in the form of X-ray fluorescence.

Each element has a characteristic emission energy for each electron shell,

referred to as K, L, and M spectra (K-alpha, K-beta, etc.), indicating from which electron shell the fluorescence originates. Figure 1 illustrates these phenomena by showing the characteristic K, L, and M spectra and their corresponding energies (in kiloelectron volts, keV).

An X-ray fluorescence spectrometer, then, is used to record these characteristic spectra and their energies, thereby identifying and possibly quantifying the elements. Four functions, therefore, were required in the PXRFS to accomplish this: (1) generate X-rays, (2) transform fluorescent energies into electrical impulses, (3) process, interpret, and store these electrical impulses, and (4) display the interpreted data in a form the operator could use.

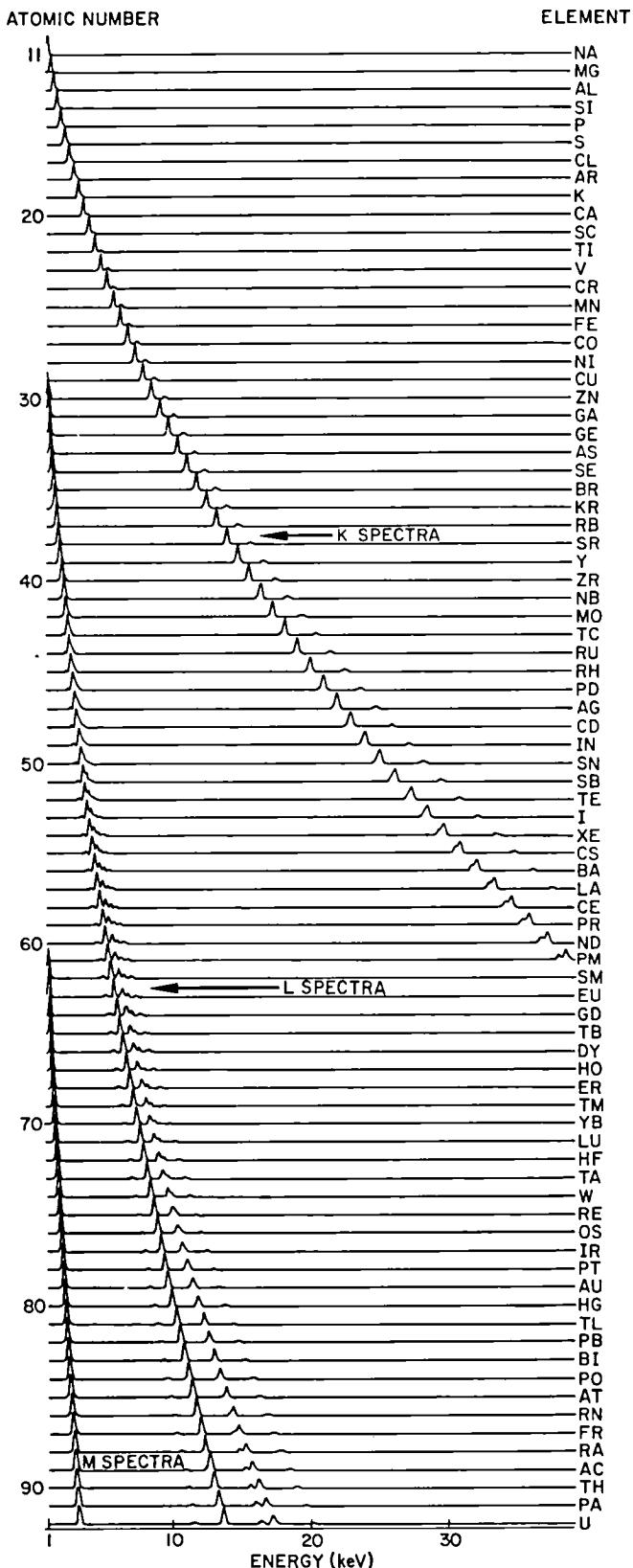


FIGURE 1. - Characteristics of X-ray fluorescence spectra. (Copyright, ASTM, 1916 Race Street, Philadelphia, Pa. 19103. Reprinted/adapted, with permission.)

INSTRUMENT DESIGN AND DESCRIPTION

Design objectives were to obtain an optimum combination of available technology and commercially available components, considering the factors of weight, size, operator training, speed of analysis, accuracy, versatility, and durability under field conditions. Within this philosophy, the optimum spectrometer design became a unit containing two X-ray sources (radioisotopes), in order to excite X-ray response from a wide range of elements, coupled with gas-proportional counters (detectors) that would feed electrical impulses (transformed fluorescent energies) to a microprocessor for processing, interpretation, and storage in memory; in turn, the microprocessor would feed data to a cathode ray tube (CRT) display and a liquid crystal display (LCD).

The PXRFS was designed for operators trained in mineral identification but lacking extensive technical knowledge of spectroscopy. The operator needs considerable experience in using the PXRFS before he/she fully understands the displays.

The PXRFS (fig. 2) consists of an analyzer unit, including a microprocessor, solid-state circuitry, batteries (power source ranges from 7 to 40 volts DC) and a control panel with a 1.3- by 1-inch CRT display, a 0.75- by 3.75-inch alphanumeric LCD, and a variety of switches and dials (described in appendix A), all connected to a sensor head by a 6-foot, spring-coiled cable. The sensor head contains two radioisotopes, 100 milli-curries (mCi) of ^{55}Fe having a calibration target composed of sulfur and titanium, and 15 mCi of ^{109}Cd having a calibration target composed of titanium and zirconium, two collimators, and two gas-proportional counters. When its cable is unplugged, the sensor head can be stored in the lid of the analyzer unit.

Dimensions of the PXRFS, lid closed, are about 11 by 12 by 5 inches (sensor head is 10.5 by 2.5 by 1.5 inches). The complete spectrometer weighs about

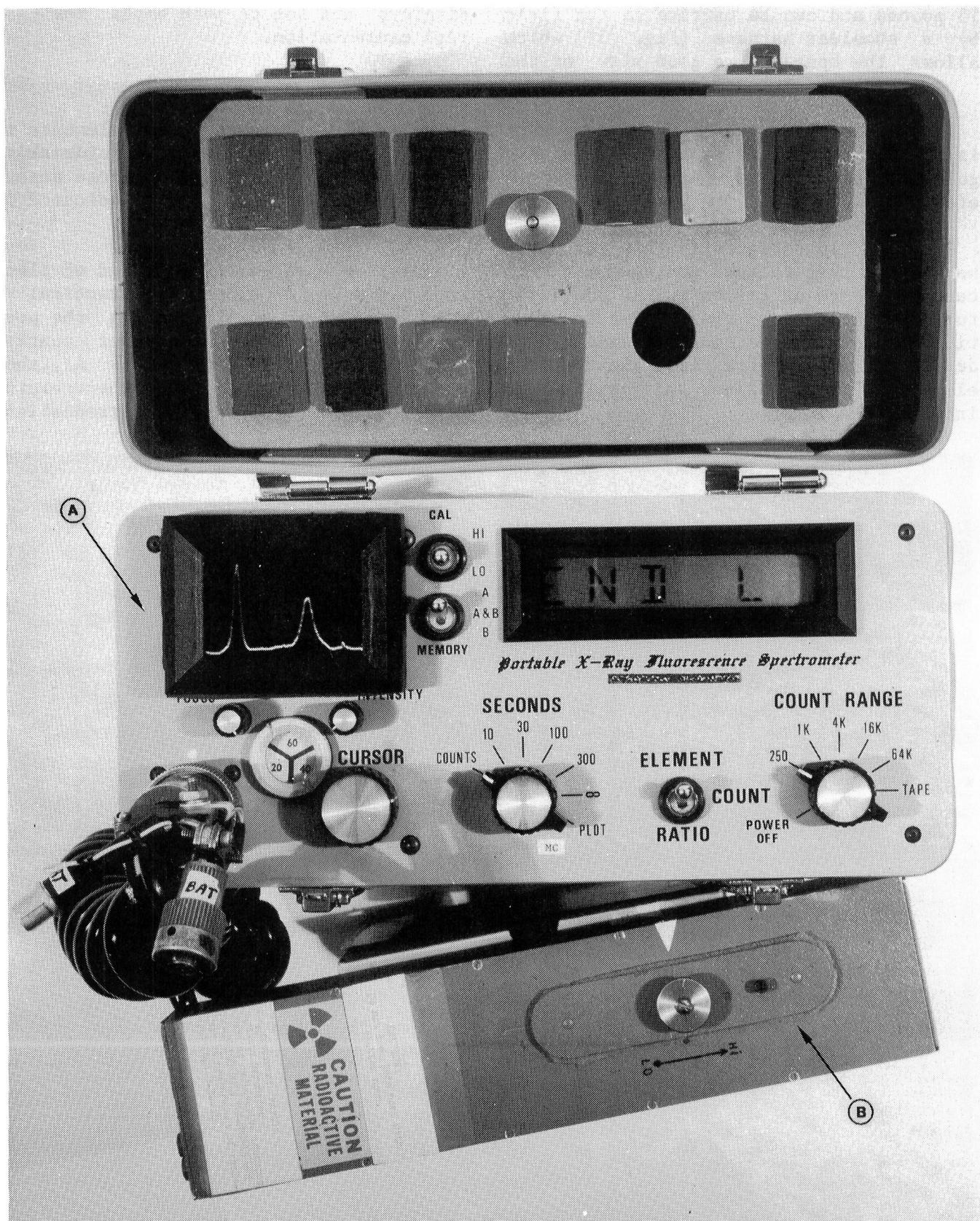


FIGURE 2. - Portable X-ray fluorescence spectrometer (PXRFS). A, Main analyzer unit; B, sensor head.

15 pounds and can be carried in the field by a shoulder harness (fig. 3) which allows the operator a good view of the

displays and use of both hands for control manipulation.

ANALYTICAL PROCEDURES AND RESULTS

Information gathered during the tests is preliminary and is presented here as a guide to prospective portable spectrometer users.

Previous investigators (bibliography) have shown that factors having a significant influence on response include moisture, particle size, the geometric relationship between the source, sample, and detector, and effects from surrounding elements ("matrix effect and interelement interference").

The useful response from elements in certain combinations varied considerably; consequently, the success of these determinations varied widely, as indicated in table B-1.

Analytical procedure consisted of placing a sample as close as practical to the sensor-head port, opening the port by sliding the port-shutter control knob (as described in appendix A), thus irradiating the sample; then the operator recorded the lapsed time of irradiation,



FIGURE 3. - PXRFS as used in the field.

total counts, elements detected, and the ratio values. The operator determines the length of analysis time by either presetting the time on the control panel or releasing the port-shutter control knob when a significant spectrum is displayed on the CRT. The spectrum is then studied and sometimes photographed by the operator.

The ratio value is derived by the microprocessor, and is a ratio between the fluorescent intensity (counts) of the element being measured and the back-scatter (BS) peak intensity, a process intended to compensate automatically for sample grain size and the geometry of the source, sample, and detector. The BS peak is created by radioisotope X-rays that escape the fluorescence process, that is, reflected X-rays recorded by the detector and displayed as a peak at the high-energy end of the spectrum.

Laboratory Testing

Theoretically, 73 elements can be detected by the PXRFS tested by the Bureau (fig. 4A). However, as of this writing, only 33 elements have been detected, as shown in figure 4B. This is attributable in part to a lack of specimens or standards that contain the other elements. Moreover, the precious metals were not detected, owing partly to the lack of spectrometer sensitivity to low concentrations of precious metals normally found in natural occurrences and partly because the ^{109}Cd isotope decays

to silver and does not excite fluorescence from silver atoms.

The PXRFS response to fluorescence (secondary X-ray emissions) of elements was investigated on 52 samples including 7 control standards loaned by Harlan Barton of the U.S. Geological Survey (USGS), Denver, Colo., 13 Bureau mineral display specimens, and 32 pulverized samples previously analyzed for a Bureau Indian-lands project (see tables B-2 through B-4).

Pulverized samples were poured into plastic 35-mm-diameter by 10-mm-deep petri dishes. Each dish was filled to heaping, tamped lightly for compaction, and leveled with a knife or straightedge, which provided a smooth sample surface for X-ray analysis. The petri dish was placed in a sample receptacle in the base of a sensor-head rack (fig. 5), which has corner uprights designed to support the sensor head at an optimum analysis distance and position over the pulverized sample.

Pulverized samples in standard kraft paper envelopes were analyzed without removal from the envelopes. The response from the ^{55}Fe source was not usable because the envelope blocked the secondary, or perhaps, the primary X-rays; response from the ^{109}Cd source appeared normal. This rapid method of analysis may effectively identify the composition of pulverized samples without removing them from the envelopes.

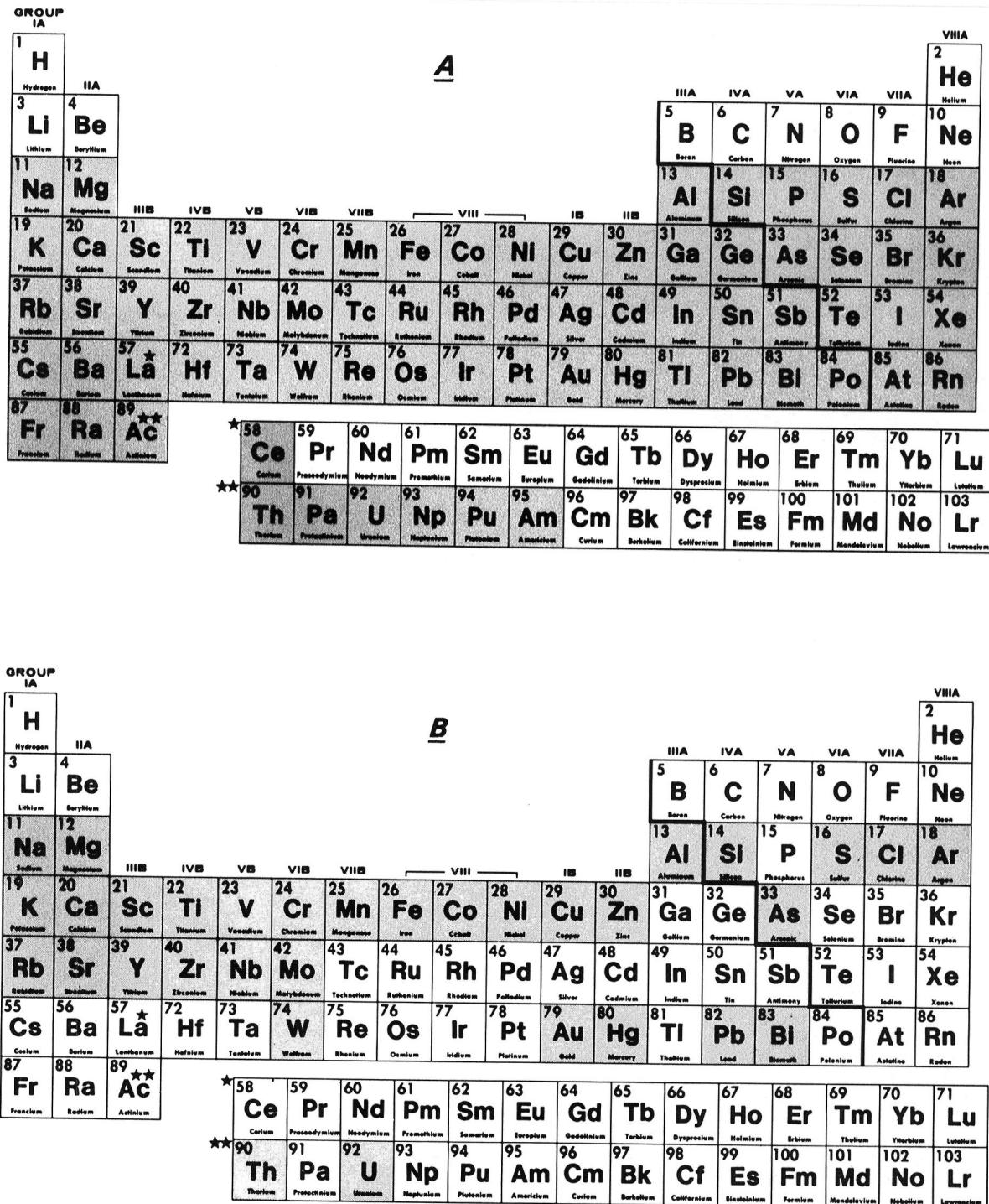


FIGURE 4. - Periodic table of elements. Shaded areas show A, elements theoretically detectable by PXRFS; B, elements actually detected in Bureau of Mines tests.

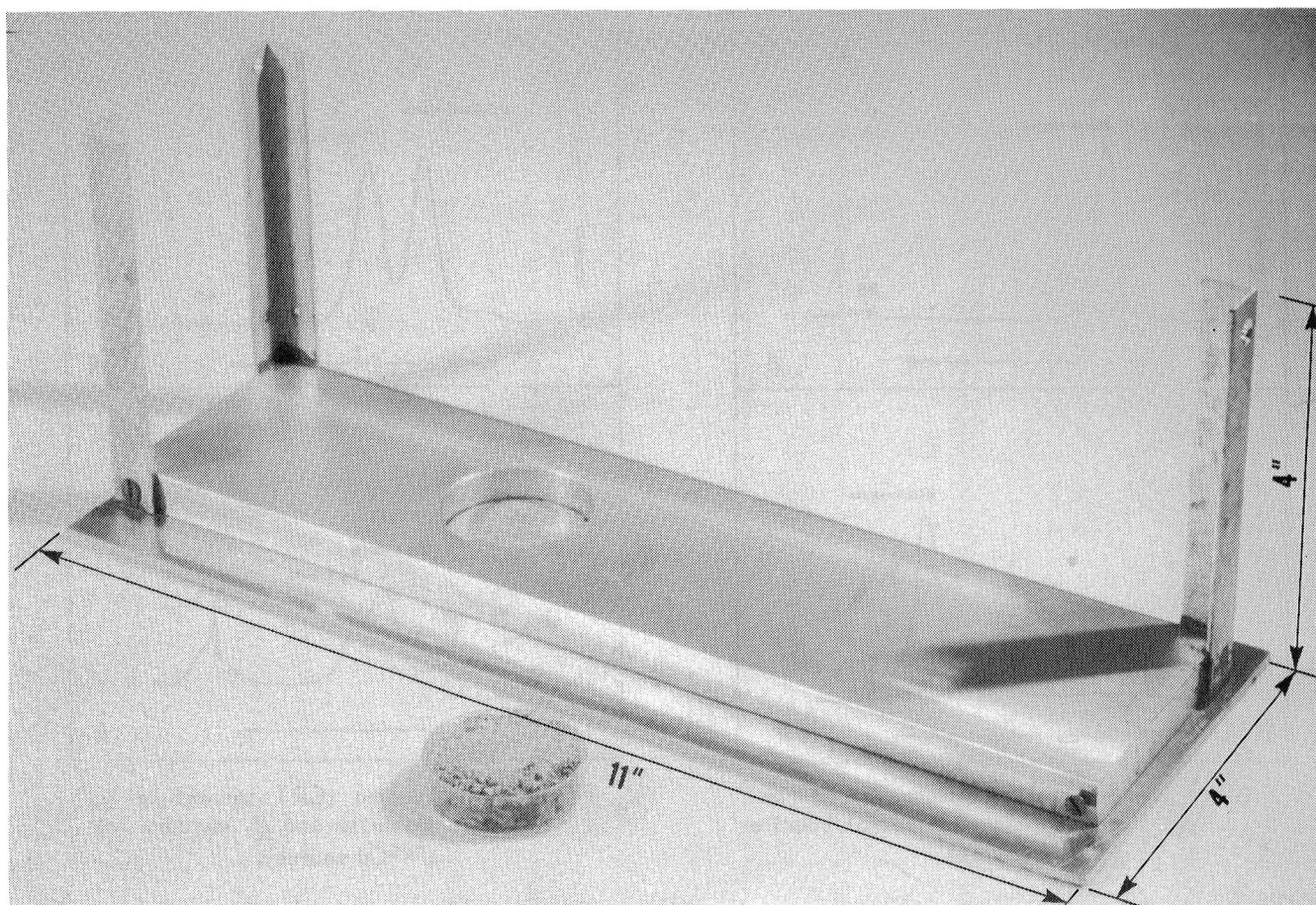


FIGURE 5. - Sensor-head rack and pulverized sample in petri dish.

Figures 6 through 9 are spectral traces from photographs of the CRT display. Each figure shows the response of certain elements or minerals. These spectral traces illustrate the fluorescent intensity (counts) plotted in relation to the element emission energies as interpreted by the PXRFS microprocessor. Shown are the element X-ray emission peaks and their spectral line classifications and the position of element emissions relative to backscatter emissions.

Interference

Because X-ray emission energies (kilo-electron volts) of many elements (fig. 1) are close in value, and because the PXRFS uses a proportional counter as a detector, fine resolution of the element emission energies was difficult. The problem was interference between elements, recorded as distortions of spectra.

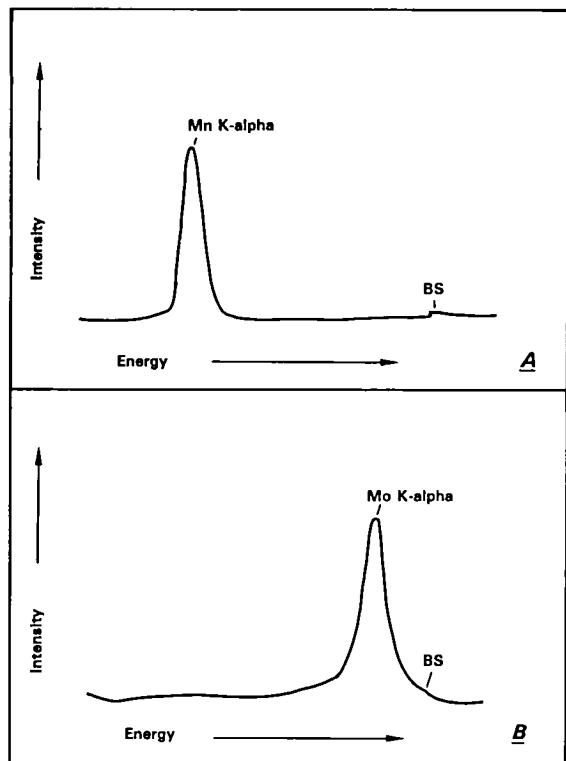


FIGURE 6. - Spectra (CRT traces) of A, manganese ore and B, molybdenite (^{109}Cd source).

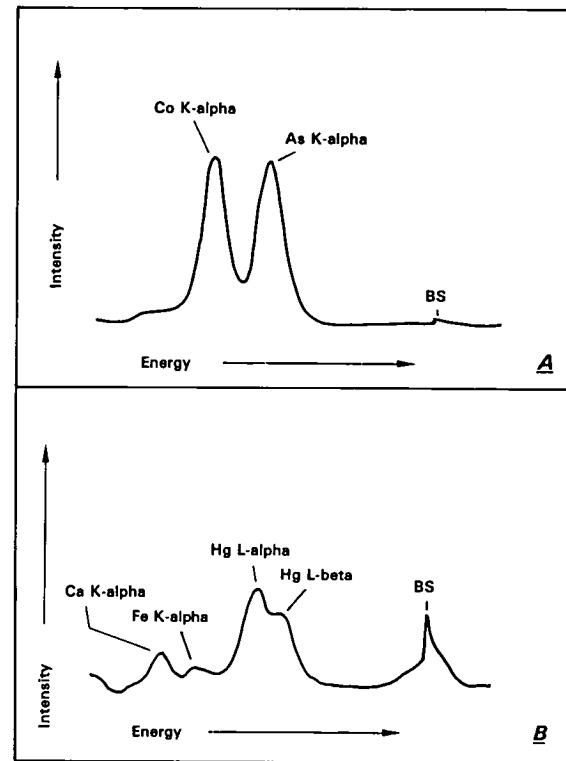


FIGURE 7. - Spectra (CRT traces) of A, erythrite and B, mercury ore (^{109}Cd source).

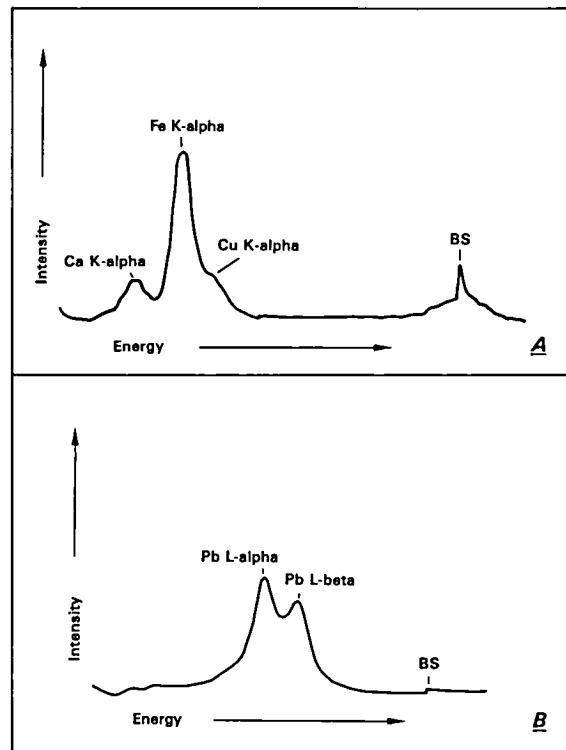


FIGURE 8. - Spectra (CRT traces) of A, sample containing iron, copper, and calcium, and B, galena (^{109}Cd source).

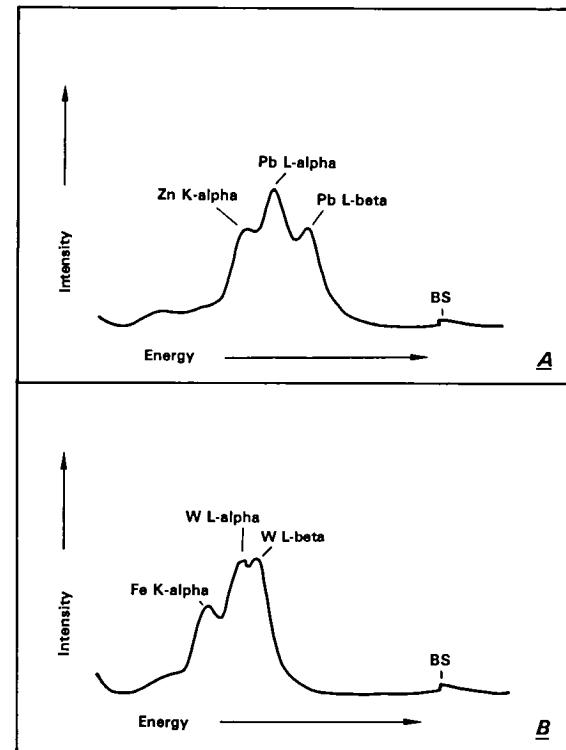


FIGURE 9. - Spectra (CRT traces) of A, sample containing lead and zinc and B, ferberite (^{109}Cd source).

Figure 10A is a spectral trace from a pulverized sample of a quartz vein in granite, which chemical analysis indicated to contain 23.7% As, 5.2% Cu, and 3.05% Pb. This spectral trace illustrates the interference created when copper, iron, and arsenic occur together. The copper emissions were interpreted as having an additive effect to the area of the spectrum between the iron and arsenic emission peaks. Also, the arsenic emissions appear to mask the lead responses; however, lead emissions have an additive effect on the arsenic emission peak. Figure 10B, a spectral trace of a pulverized sample of galena in an igneous rock, which contains 1.32% Pb and 1.95% Zn, illustrates further the interference problem by showing the distortion of the predominant iron emission peak by the lead and zinc emissions.

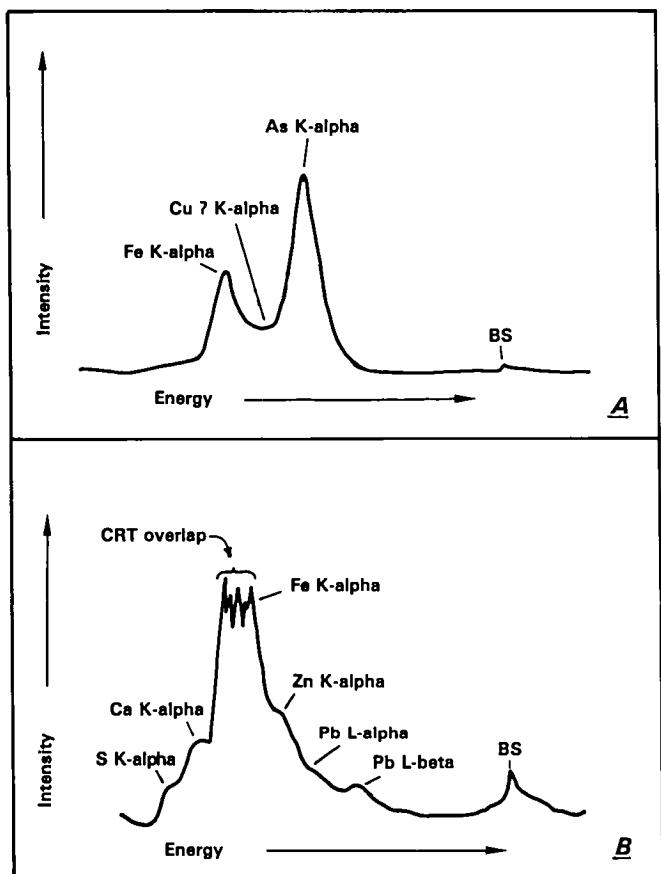


FIGURE 10. - Spectra (CRT traces) illustrating inter-element interference. A, Sample containing As, Cu, Pb, and Fe; B, high-Fe sample containing Zn and Pb (^{109}Cd source).

Iron was the most significant source of interference, owing to its natural abundance and strong X-ray emission response. Many elements of economic interest occur with iron; because iron interfered with the X-ray responses of these elements, iron interference became the major problem. In tests on certain clays (ceramic raw materials) even minor amounts of iron impurities were detected; the PXRFS could be used for the selection of low-iron clays for industrial purposes.

With experience, the PXRFS operator was able to recognize characteristic emission peaks of elements even though the peaks were modified by interference.

Matrix Effect

The matrix effect is a form of interference simply defined as the effect matrix elements have on the X-ray response of other elements. Matrix effect is due to an interaction between the emission and absorption characteristics of the matrix elements and the emission energies (or wavelengths) of the elements being analyzed.

Rock-forming or gangue minerals (matrix) can have a significant effect on the X-ray responses from elements of possible economic interest. If rock composition changes, the matrix effect also changes. That is, as the relative proportion of matrix elements changes, their effect on economic elements changes in the form of an increase or decrease in secondary radiations or emission absorptions.

Summary of Results

Positive identification and semiquantitative analyses of elements were possible for many of the samples tested. The limitations created by the detection system used in the PXRFS, however, precluded utilization of the instrument's full potential.

Element identification was possible for most of the elements tested and was

affected only by the presence of interfering elements for element concentrations less than 0.10%.

Quantitative analyses were hindered by both interelement interference and matrix effect, the matrix effect especially affecting the results when element concentrations were low (generally less than 0.10%). Semiquantitative results probably can be obtained by use of the PXRFS, but it will be necessary to use samples (standards) that have known but varying percentages of element concentration for comparison with samples of unknown composition. Five element calibration curves (fig. 11), constructed from USGS standardized samples, are plots of element concentrations relative to the ratio values calculated by the PXRFS. Quantification of elements in a sample of unknown composition can be estimated by using calibration curves if the composition of the sample approximates that of the standards used to make the calibration curves.

The overall quantification results, based on spectral emission data as compared to analyses of the samples, varied from useful to nearly useless, depending on the relative percentages of interfering elements as well as the X-ray response characteristics of a particular element and those elements adjacent to it in the spectrum.

Field Testing

On short field trips to selected mining districts west of Denver, Colo., and on a field trip to southwestern Arizona, the PXRFS was tested for its durability and its practicability in addition to its analytical accuracy and response capabilities.

The PXRFS was transported in a four-wheel-drive vehicle to selected test sites over varying types of roads. During transportation, the instrument was carried in a high-impact plastic case (fig. 12).

Hand specimens collected from outcrops or mine dumps and outcrop faces were

analyzed during the field tests (fig. 13). Some hand specimens and outcrop faces were difficult to analyze with the PXRFS because of their surface irregularities. One-handed operation of the sensor head during field tests while climbing on rock outcrops proved to be difficult because of the sensor head shape and the tension on the port shutter control return spring.

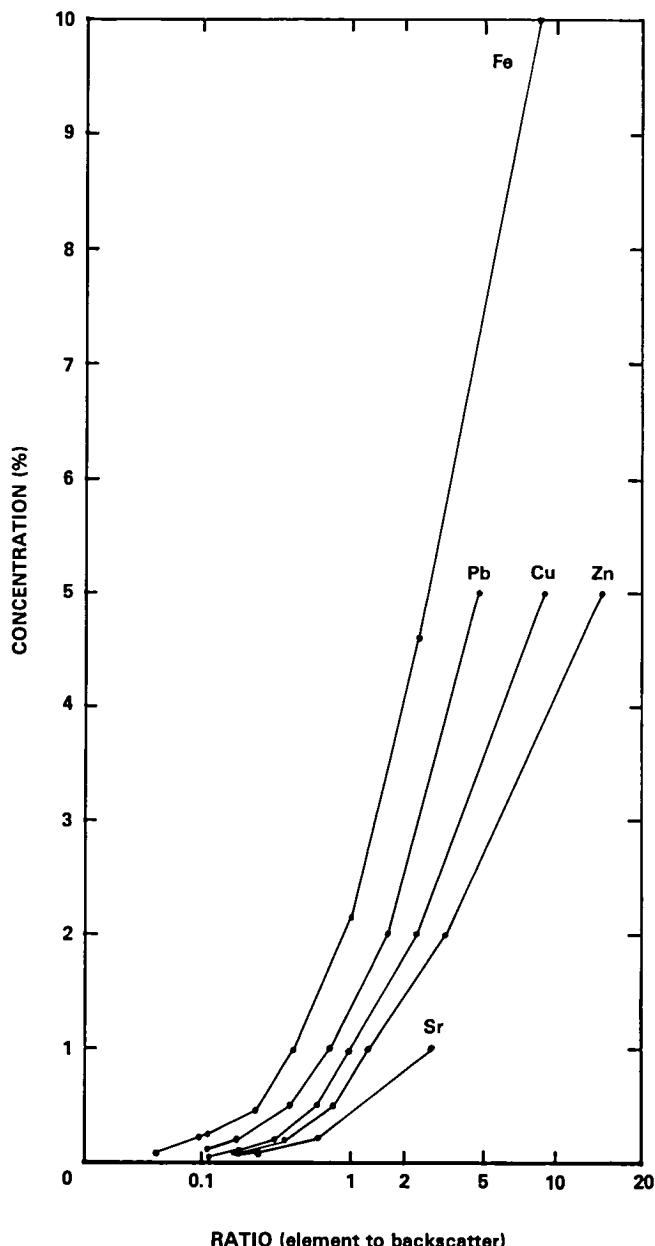


FIGURE 11.- Calibration curves of five element standards having granitic matrixes.



FIGURE 12. - High-impact plastic transport case for spectrometer.

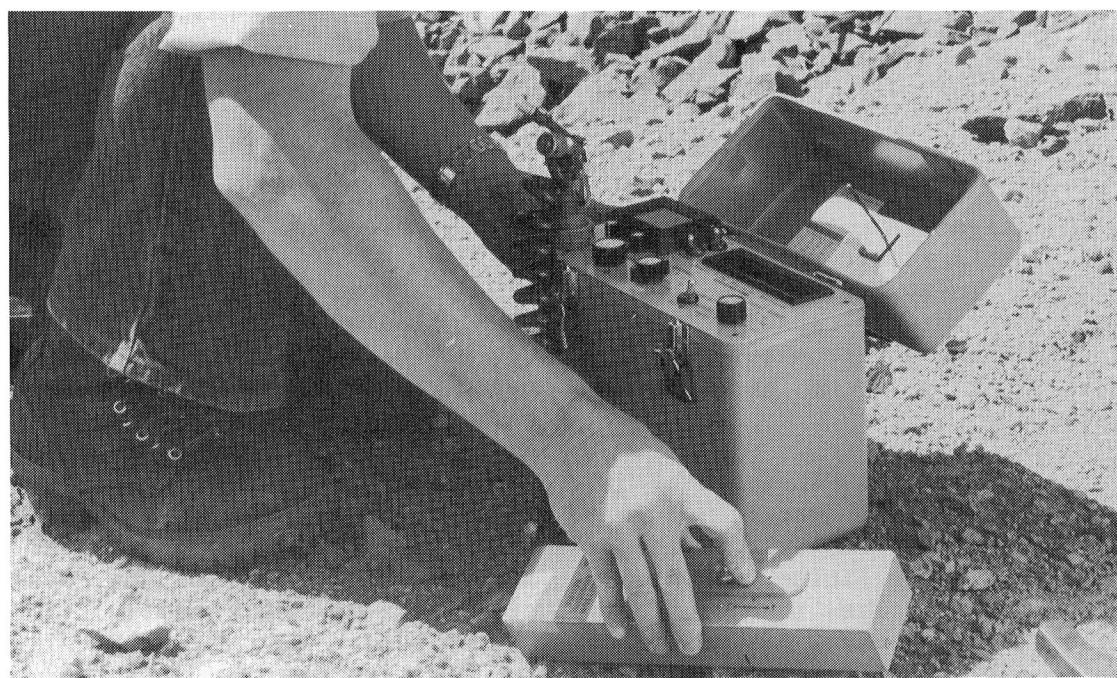


FIGURE 13. - Spectrometer in use on a mine dump.

The PXRFS response was compared with visual, physical, and chemical mineral-identification techniques; siderite, sphalerite, ferberite, and powellite were among the minerals identified during the field tests.

The tests disclosed that air temperatures between 50° and 90° F had little effect on the PXRFS; however, the instrument did undergo a spectral shift toward the low-energy end of the spectrum at 108° F. Figure 14A shows the ^{109}Cd calibration spectrum (trace) produced by the PXRFS within the operating temperature range; figure 14B illustrates the spectral shift that occurs to the same calibration spectrum at 108° F. At air temperatures less than 50° F, the spectrum appeared to shift slightly toward the high-energy end of the spectrum. The spectral shift phenomenon made element identification difficult and necessitated calibration adjustments in the sensor head that were time consuming and not always successful.

Durability of the PXRFS was somewhat less than expected. A fuse holder attached to the chassis broke loose; wires in the sensor head broke twice from flexure; a beryllium window in the ^{109}Cd -coupled detector shattered, which made the detector and the PXRFS inoperable;

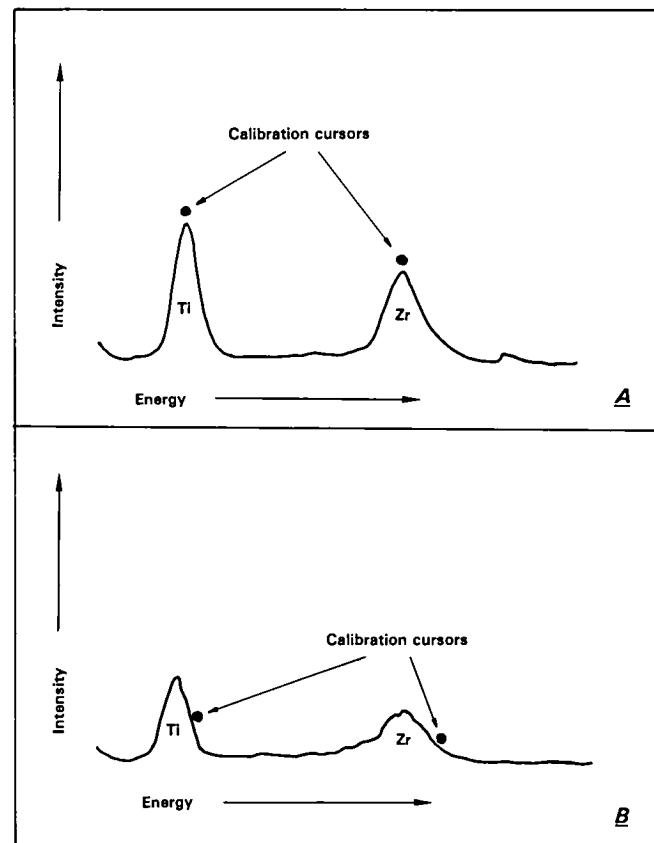


FIGURE 14. - Calibration spectrum of ^{109}Cd source.
 A, under normal operating temperature;
 B, at 108° F.

and the analyzer unit lid hinges were sprung from the weight of the sensor head.

CONCLUSIONS

This project involved limited testing of the PXRFS for field use in mineral-resource investigative work, and results presented in this report should be considered preliminary. Testing demonstrated a portion of the capabilities and limitations of the spectrometer in mineral identifications and quantitative determinations.

The advantages of the PXRFS are (1) it is portable and can be carried in the field with little difficulty, (2) it has repeat analysis capabilities, (3) analysis time is generally less than 1 minute (if the radioisotopes have not undergone a decay of one half-life), (4) all data are available instantly on the displays,

(5) no sample preparation is necessary in the field, and (6) it is relatively maintenance free.

Limitations of the PXRFS are (1) poor resolution of emission data owing to use of a gas-proportional detector that causes interference problems, (2) short half-life ($^{109}\text{Cd} = 1.2$ years, and $^{55}\text{Fe} = 2.6$ years) of the radioisotopes results in periodic replacement expense, (3) durability is somewhat less than expected, (4) the calibration adjustment screws, located in one end of the sensor head, are relatively inaccessible, (5) spectral shifts occur when the PXRFS is operated in air temperatures less than 50° F or more than 90° F, (6) the CRT display is

hard to read because of its miniature size, or because of bright daylight in the field, and (7) one-handed operation of the sensor head is difficult because of its shape and port shutter control spring tension.

At this time the PXRFS falls short in achieving the desired objective of reliable quantitative analytical results.

RECOMMENDATIONS

Recommendations for modification of the PXRFS mainly involve the detectors and the radioactive sources. In order to improve spectral resolution and reduce operating expenses of the PXRFS, the following modifications are suggested:

1. Replace the gas-proportional counters with a single mercuric iodide detector to enhance spectral resolution.
2. Replace the ^{109}Cd and ^{55}Fe radioisotopes with an americium (^{241}Am) source, thereby increasing the element excitation range and reducing isotope replacement costs (the ^{241}Am source has a half-life of 458 years).

A miniature X-ray generator may be an alternative to radioactive isotopes as an X-radiation source.

Other modifications to the PXRFS, for special-use purposes (such as ore control sampling) and to further reduce cost and complexity, might include the following: Use only one radioactive source; test a weaker radioactive source (at lower cost); eliminate the CRT display; use a three- or four-digit LCD and a manual element selector (cursor control in present PXRFS) with a meter (analog) readout.

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APPENDIX A.--SELECTED SECTIONS OF PXRFS OPERATION
AND MAINTENANCE MANUAL¹

Figure A-1 provides a display of the front panel and controls of the PXRFS as a reference for the instructions contained in this section.

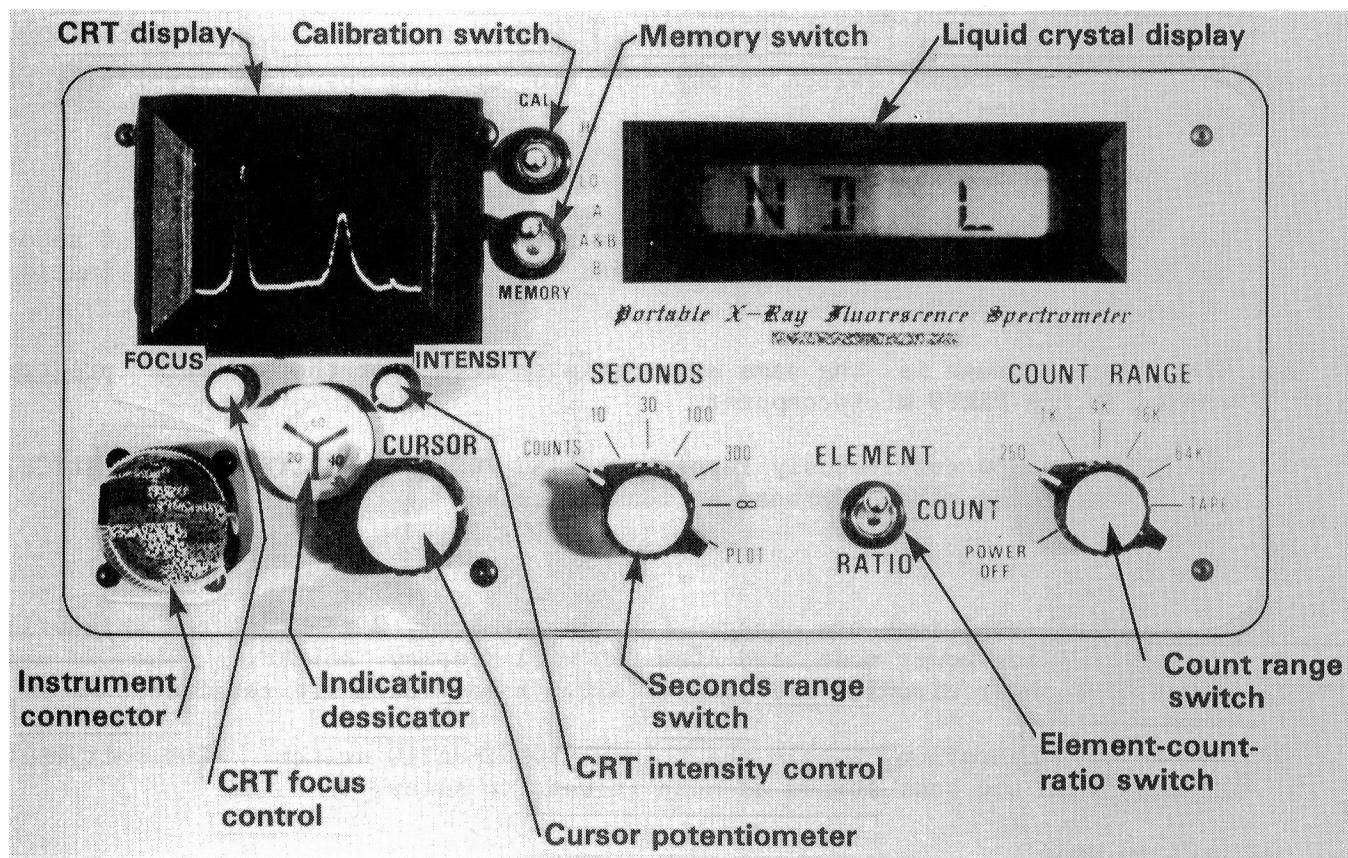


FIGURE A-1. - Front panel and controls of PXRFS.

¹Martin Marietta Corp., rev. July 11, 1980.

OPERATING INSTRUCTIONS

1. Power on

- A. Connect sensor head to instrument by pushing the sensor head cable connector onto the face panel connector (aline before plugging in) and then turning clockwise for approximately one-third turn.
- B. Turn the COUNT RANGE switch to any of its numbered positions (typically to the 250 position).
- C. Set the SECONDS switch to 30.
- D. The liquid crystal display (LCD) shows the results of the instrument automatic self-test. It will display either "SELF TEST PASSED" or a combination of warning messages.
- E. The next message is the name and revision level of the software program stored in the PXRFS microcomputer.
- F. The LCD will now continuously repeat "WAITING FOR OPEN SHUTTER OR CAL SWITCH" until one or the other of these actions is taken.

2. Sleep Mode

- A. If no switch positions are changed for approximately 2 minutes, the unit will switch to a low-power mode, and the LCD will display "ASLEEP." The CRT and other high power circuits are powered off. Memory data are retained.
- B. To resume operation, toggle the ELEMENT-COUNT-RATIO switch. The unit will wake up with the same status as when it went to sleep.

3. Calibration Mode

- A. Calibration should be performed before starting a new series of analyses.
- B. After the power-on step, with the LCD repeating the "WAITING..." message, set the SECONDS switch to either COUNTS or to one of the numbered seconds positions.
- C. Set COUNT RANGE switch to desired scale (typically 250 for calibration).
- D. Verify sensor head shutter is closed.
- E. Toggle CAL switch momentarily to L0. LCD displays "Fe-55" for 2 seconds, then begins displaying elapsed analysis time as the spectrum is accumulated.
- F. The unit is now analyzing the Fe-55 calibration sample (sulfur and titanium target) built into the sensor head.
- G. Adjust CRT focus and intensity knobs for best viewing of the spectrum.
- H. When analysis is complete the LCD will display "CAL," and the CRT display will be continuous with two flashing cursors. The left cursor marks where the sulfur peak should appear, and the right cursor where the titanium peak should appear.

NOTE: At any time during or after analysis the COUNT RANGE switch may be used as desired to adjust the vertical expansion of the spectrum on the CRT display.

- I. If the cursors are not on the two peaks, adjust the LO trimpot on the sensor head and repeat the analysis as necessary. After adjusting the potentiometer, restart the spectrum by operating the CAL switch again.
- J. After the Fe-55 channel is calibrated, repeat the above procedure for Cd-109. Substitute CAL HI for CAL LO and use the HI trimpot on the sensor head. For Cd-109 the left cursor marks titanium and the right cursor marks zirconium.

4. Normal Operation

- A. After power-on and calibration steps, the unit is ready to analyze samples.
- B. To analyze, adjust the SECONDS and COUNT RANGE switches for the desired type of analysis. Set the MEMORY switch to either A or B, depending upon which half of memory the spectrum is to be stored. Use the A and B setting only if there is no need to compare two different spectra.
- C. Place the sensor head port against the sample of interest. If this is not possible, place the port as close to the sample as possible (see "Safety" section at the end of the appendix).
- D. Using an index finger, press and hold the shutter control either forward for a Cd-109 analysis, or backward for an Fe-55 analysis. The LCD will first identify which radiation source is in use and then the elapsed analysis time (in seconds). The beeper in the sensor head will sound to warn that a radioactive source is exposed through the open shutter.

NOTE: If the beeper doesn't sound when the shutter is opened, notify the instrument manufacturer.

- E. While waiting for the analysis to complete, observe the spectrum building up on the CRT. When sufficient data have been taken, release the shutter control. Verify that the alignment circle is properly centered in the port window, indicating the spring-loaded shutter is in the fully closed position.
- F. The COUNT RANGE switch may be used to expand the CRT display as desired.

CAUTION: Do not turn the COUNT RANGE switch to the 9 o'clock position, or the unit will be powered off and special data lost.)

- G. To identify a peak, use the CURSOR knob to adjust the bright dot to the top of the peak of interest. When the cursor is adjusted to an element channel, it will flash once per second, and with the ELEMENT-COUNT-RATIO switch set to ELEMENT, the LCD will display the element's chemical symbol.
- H. For a hard-copy record of the spectrum, connect the plot or tape output wires to a chart recorder or tape recorder, respectively. These wires are available from the connector at the front panel end of the sensor cable.

1. For a plot, set the SECONDS switch to PLOT. The LCD will display "PLOT REQUESTED" and "PLOT." When the "PLOT" message appears the unit will begin outputting 0- to -5-volt analog signals of spectral data at eight channels per second. The spectrum will repeat as long as the SECONDS switch is left on plot. Switching off the PLOT position at any time will terminate the output and return the spectrometer to the previous mode.
2. For recording the spectrum onto magnetic tape, connect the tape output plug to the auxiliary or microphone input of a recorder.

NOTE: It may be necessary to adjust the drive circuit output impedance to match the recorder input impedance. Turning the TAPE trimpot, on the INTERFACE printed circuit board, clockwise will increase the impedance to 60 kilohms maximum; turning counter-clockwise will decrease it to its 10-kilohm minimum.

Set the COUNT RANGE switch to TAPE and then put the recorded in the record mode. The LCD will display "TAPE DUMP REQUESTED" and "RECORD" (signals start of dump). The spectrum will be dumped only once, and the dump is not interruptable (takes approximately 2 minutes).

- I. To perform more analyses, either leave the MEMORY switch in the same position it was, or change from A to B or from B to A to save the last spectrum taken. Repeat the steps under section 4.

5. Charging the Batteries

- A. Be sure the COUNT RANGE switch is in the power-off position.
- B. Unplug sensor head cable by twisting counter-clockwise for approximately one-third turn and pulling to disengage.
- C. Plug in cable from the charging unit to the front panel connector.
- D. Place the CHARGE switch on the charging unit to one of its three positions. Under standard conditions, the unit contains nickel-cadmium batteries, but lead-acid may also be installed. Use LO RATE for nickel-cadmium batteries if the time available for charging is 16 hours or more; if a fast charge is needed, or if the unit is to be operated during charging, use the HI RATE setting.
- E. Connect the appropriate terminals to either the 110-volt-ac source, or automotive battery (12 volts dc).

ROUTINE MAINTENANCE

1. Changing Sensor Heads

- A. Place the COUNT RANGE switch in the power-off position.
- B. Unplug sensor head at the front panel connector of the main unit by twisting counter-clockwise for one-third turn and pulling off.
- C. Plug in new sensor by alining connector, pushing to mate pins, and rotating connector clockwise to snap into locking position.

2. Changing the Desiccant Tube

- A. Replace or rejuvenate desiccant whenever the 60% sector changes from its normal blue color to either pale pink or white.
- B. Remove the cartridge from the front panel by unscrewing it with fingers or a suitable straightedge.
- C. Screw in new unit, or rejuvenate removed cartridge by baking in an oven at 125° C (257° F) for 2 hours or more.

3. Replacing the Batteries

The unit is powered by seven D-size batteries. Either rechargeable batteries (for example, nickel-cadmium or lead-acid cells) or nonrechargeable cells (for example, carbon-zinc, mercury, or lithium types) may be used. Standard rechargeable batteries are of the nickel-cadmium type, but the lead-acid type is preferred for operation in very cold weather.

- A. Turn power off and unplug the sensor head. Remove the hinged lid (optional).
- B. Unscrew and remove the four mounting feet at the base of the instrument.
- C. Carefully place the unit on the side where the four mounts were removed (front panel facing up).
- D. Lift up on the front panel to pull the unit slowly out of the housing (assistance from a second person will be helpful and is a sensible precaution).
- E. Set the unit down on a smooth surface.
- F. Note the battery orientations. All cells are aligned in the same direction, with negative terminals closer to the CRT than the positive terminals.
- G. Remove batteries from holders by pushing end clips toward each other, and releasing tiedown clip with a slight twisting motion.
- H. Place new battery in proper direction and again press end clips inward to facilitate installing the tiedown clip in its original orientation.
- I. Replace one or more batteries as required.
- J. Carefully place unit in housing, align holes at bottom with ports, and screw in mounting feet.

TROUBLESHOOTING

The first thing to suspect in the event of improper operation of the unit is a weak or dead battery. Even when the unit is powered by an external electrical source, a malfunctioning cell can prevent proper operation. Therefore, first try changing the batteries (see step 3 of the "Routine Maintenance" section) to verify that is not the problem. During this operation, check for any obvious internal problems: a printed circuit board that is not seated into its connector, a broken wire, dirt or other contamination in critical locations, charred or burned spots, etc.

If this does not solve the problem, and the unit is known to have been exposed to high humidity or low temperatures, remove the desiccant tube and place the entire unit in a dry, warm environment to remove condensation. Replace the tube with a fresh or rejuvenated desiccant.

In the case that the calibration peaks do not line up properly and cannot be adjusted into position with the appropriate potentiometer, the unit will require resetting by the manufacturer. However, it is possible to make temporary use of the instrument for field purposes by using the calibration targets built into the lid to note the channel numbers into which the elements of interest fall. With this approach, the operator must disregard the element symbols given on the LCD since these peaks will now occur slightly upscale or downscale from the normal position.

SAFETY

The PXRFS instrument has been engineered with every consideration for safety. A number of special features are designed to minimize the possible risks to the operator and bystanders. It is extremely important, however, that all persons who are using, or have plans to use this instrument, read this section very carefully. THIS UNIT CONTAINS RADIOACTIVE MATERIALS and although these sources emit relatively low energy X-rays, and are carefully shielded, there is the possibility of unnecessary radiation exposure whenever the shutter is opened without the sensor head being placed flush against a suitable thick sample.

1. Radiation Hazard

The standard sensor head contains two radioactive materials: approximately 100 mCi of iron-55 isotope and 15 mCi of cadmium-109. These sources are electroplated onto a substrate and then hermetically sealed into a rugged holder. The possibility of leakage of radioactive material is extremely remote. The sources are in turn mounted in shielded collimators. When in the neutral closed position the safety shutter provides additional shielding to prevent any radiation from penetrating outside the sensor head. There is no hazard associated with handling or being near the sensor head when the shutter is closed (that is, when neither source is exposed). Of course, the sensor head should be kept secured from access by casual bystanders at all times.

During acquisition of a spectrum, one or the other of the two sources must necessarily be positioned to irradiate the samples with X-rays. To prevent unnecessary exposure to the operator and for maximum safety, the following rules must be followed:

- A. Always place the sensor head analysis port up against the sample before opening the shutter.
- B. If the sample surface is irregular or it is impractical to place the sensor head totally against the sample, be sure to position your hands, limbs, and body such that they are behind the sensor head.
- C. Never look at or handle the port window, except when the shutter is fully closed.
- D. If the warning beeper fails to operate when the shutter is opened, return the unit to the manufacturer.

- E. Never attempt to repair or disassemble the sensor head. The only exception is simple replacement of the plastic film port window using a pre-prepared window kit. Make sure the shutter is in the neutral closed position before making this change.
- F. Insure that the radiation sources are leak checked at required intervals (typically, every 6 months).
- G. Have available a thin-window Geiger counter or other radiation monitoring device. Periodically check the radiation level during normal operation to make sure unsafe procedures are not being followed.

2. Shock Hazard

All circuits within the PXRFS sensor head operate at low voltages, with the exception of the detector bias supply. This supply provides 1,000 to 1,500 volts for operation of the proportional counters. Because this supply is potted, shielded by a metal housing, and contains a current-limiting resistor, the possibility of a malfunction causing serious shock is minimal. In the event of any indications of electronic problems within the sensor head, the unit should be returned to the manufacturer.

An additional high voltage circuit is located within the main unit to provide the electron beam accelerating voltage for CRT operation. Because of this supply, it is recommended the unit never be turned to power-on condition when removed from its housing--for example, during a battery change-out operation.

APPENDIX B.--TABLES OF ANALYTICAL RESULTS

TABLE B-1. - X-ray excitation capabilities of ^{55}Fe and ^{109}Cd for selected elements

Element	X-ray source used in analysis	Source excitation capabilities	Most likely interfering elements
Aluminum.....	^{55}Fe	Fair.....	Mg, Si
Arsenic.....	^{55}Fe	Good.....	Pb, Se, Hg
Calcium.....	^{55}Fe and ^{109}Cd	Fair.....	K
Chromium.....	^{55}Fe and ^{109}Cd	Good.....	Fe, V, Mn
Cobalt.....	^{109}Cd	Good.....	Mn, Fe, Ni, Cu
Copper.....	^{109}Cd	Good.....	Fe, Co, Ni, Zn
Iron.....	^{109}Cd	Good.....	Cr, Mn, Co, Ni, Cu
Lead.....	^{109}Cd	Good.....	As
Magnesium.....	^{55}Fe	Fair.....	Na, Al
Manganese.....	^{109}Cd	Good.....	Cr, Fe
Mercury.....	^{109}Cd	Fair.....	As, Pb
Molybdenum.....	^{109}Cd	Good.....	Nb, U
Nickel.....	^{109}Cd	Good.....	Cu, Co, Fe
Niobium.....	^{109}Cd	Fair.....	Th, U
Silicon.....	^{55}Fe and ^{109}Cd	Fair.....	Al
Sulfur.....	^{55}Fe and ^{109}Cd	Fair.....	K
Titanium.....	^{55}Fe and ^{109}Cd	Good.....	Ca, V
Tungsten.....	^{109}Cd	Good.....	Cu, Zn
Uranium.....	^{109}Cd	Fair.....	Mo, Th, Sr, Nb
Vanadium.....	^{55}Fe and ^{109}Cd	Fair.....	Ti, Cr, Mn
Zinc.....	^{109}Cd	Good.....	Cu, W
Zirconium.....	^{109}Cd	Good.....	Th, Nb

TABLE B-2. - Comparison of minus 200-mesh pulp sample analysis data with PXRFS test results

(Ratio value: Total counts divided by backscatter)

Ele- ment	Sample analysis, %	PXRFS results		Comments	Ele- ment	Sample analysis, %	PXRFS results		Comments
		Total counts	Ratio				Total counts	Ratio	
GALENA IN IGNEOUS ROCK									
Ca..	NA	87	2.937	ESA; 60-sec ^{109}Cd irradiation.	Ca..	NA	27	7.375	ESA; 60-sec ^{109}Cd irradiation.
Fe..	NA	1,173	46.000	The Fe response interferes	Cu..	11.00	153	30.750	Cu response interferes with
Pb..	1.32	51	1.375	with the Zn and Pb X-ray	Fe..	NA	133	35.812	the Zn response. See
S...	NA	37	1.437	response. See trace 1.	Pb..	14.00	82	18.562	trace 3.
Zn..	1.95	77	3.500		S...	NA	9	2.812	
SCHEELITE-POWELLITE IN FAULT ZONE									
Ca..	NA	186	0.312	ESA; 300-sec ^{109}Cd irradia-	Zn..	.11	ND	NAP	ESA; 80-sec ^{109}Cd irradiation.
Cu..	NA	81	.375	tion. Fe response interferes	Ca..	NA	77	4.812	See trace 4.
Fe..	NA	258	1.062	with Cu response--Cu appears	Cu..	.018	ND	NAP	
Mo..	0.0095	60	.187	as small shoulder to right of	Fe..	NA	287	14.125	
W...	.3200	77	.375	Fe peak; W plateau to right	Pb..	2.800	89	4.375	
				of Fe peak. See trace 2.	S...	NA	31	1.312	
					Ti..	NA	72	4.187	
					Zn..	23.500	1,075	52.562	

See notes at end of table.

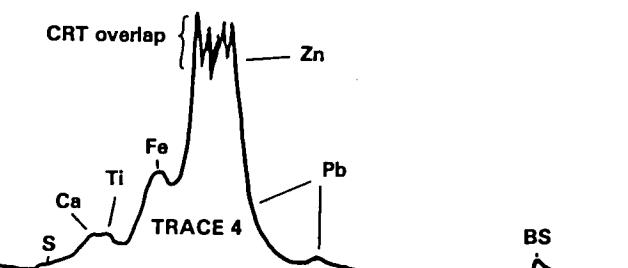
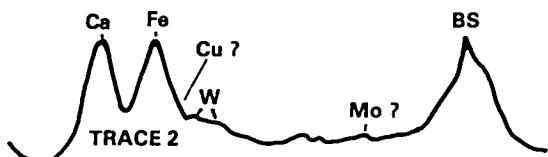
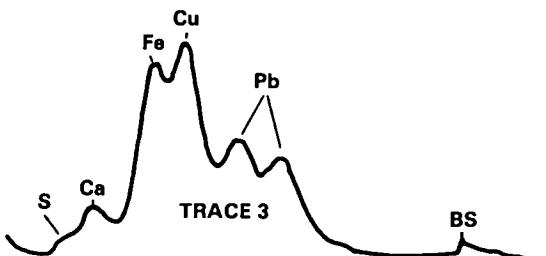
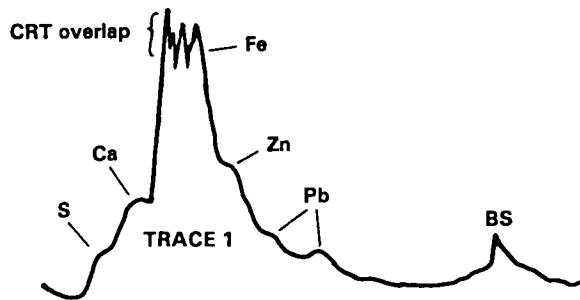


TABLE B-2. - Comparison of minus 200-mesh pulp sample analysis data with PXRFS test results--Continued

Element	Sample analysis, %	PXRFS results		Comments	Element	Sample analysis, %	PXRFS results		Comments
		Total counts	Ratio				Total counts	Ratio	
SULFIDE									
Ca..	NA	132	4.187	ESA; 60-sec ^{109}Cd irradiation.	Ba..	0.0	ND	NAp	ESA; 120-sec ^{109}Cd irradiation.
Fe..	NA	229	6.875	The Pb response is masked by Fe and Zn responses. Fe appears as a shoulder to left of Zn peak. See trace 5.	Ca..	7.0	ND	NAp	Fe appears to be shoulder on the right side of the Ti peak (trace 7). Ti peak may have intense Ca response contribution rather than just Ti response.
Pb..	0.4	ND	NAp		Fe..	2.0	41	1.562	
Zn..	NA	551	18.500		Mg..	2.0	ND	NAp	
ALTERED RHYOLITE PORPHYRY									
Ca..	NA	32	0.312	AA; 120-sec ^{109}Cd irradiation.	Pb..	.3	ND	NAp	
Cu..	0.1000	18	.187	Mo detection may indicate only a background value and ratio. Bulge in right side of Fe peak (trace 6) is a distortion caused by the Cu response.	Sr..	.4	63	2.875	
Fe..	NA	99	.750		Ti..	.03	172	7.312	
K...	NA	53	.312		Zn..	.6	45	1.812	
Mo..	.0004	18	.182		STREAM SEDIMENT FINES				
S...	NA	13	.125		Ca..	NA	37	0.375	ESA; 90-sec ^{109}Cd irradiation.
Sc..	NA	46	.312		Cu..	0.0125	ND	NAp	See trace 8.
W...	NA	28	.125		F...	.0680	ND	NAp	
Zn..	NA	12	.125		Mo..	.0330	47	.187	

See notes at end of table.

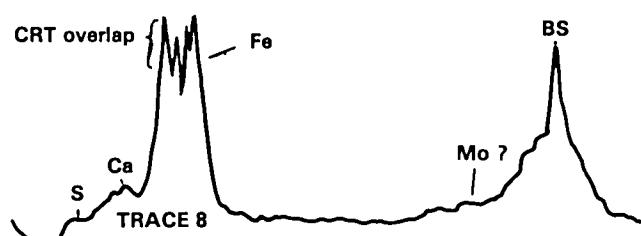
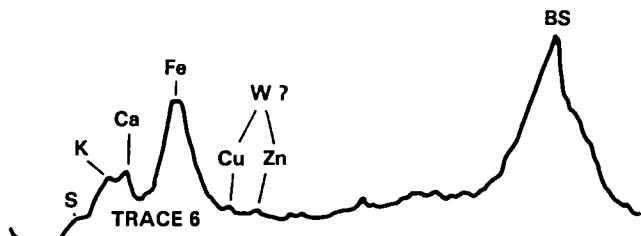
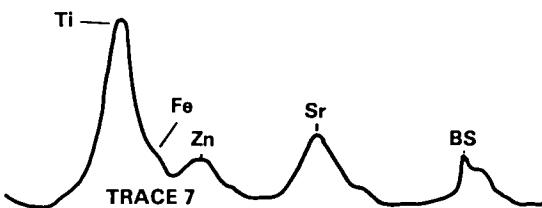
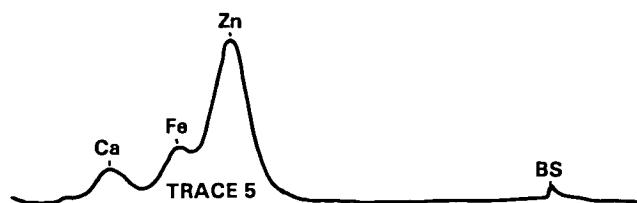


TABLE B-2. - Comparison of minus 200-mesh pulp sample analysis data with PXRFS test results--Continued

Element	Sample analysis, %	PXRFS results		Comments	Element	Sample analysis, %	PXRFS results		Comments
		Total counts	Ratio				Total counts	Ratio	
STREAM SEDIMENT FINES--Continued									
Ca..	NA	67	0.312	AA; 60-sec ^{109}Cd irradiation.					questionable; however, Cu is known to occur in this sample--Zn possibly in sphalerite as well.
Cu..	0.0025	ND	NAp	Mo interference from back-scatter peak. See trace 9.					
F...	.0520	ND	NAp						
Fe..	NA	313	1.375						
Mo..	.0480	60	.250						
W...	.0042	ND	NAp						
QUARTZ VEIN IN GRANITE--Continued									
As..	0.50	286	3.375	ESA; 120-sec ^{109}Cd irradiation. As, Bi, and Pb	As..	0.50	ND	NAp	ESA; 60-sec ^{55}Fe irradiation. See trace 11.
Bi..	.05	274	3.125	counts and ratio readings	Bi..	.05	ND	NAp	
Ca..	.10	49	.625	are all very close in value; interference among these	Ca..	.10	ND	NAp	
Cu..	.05	57	.750	three elements (trace 10) makes analysis difficult.	Cu..	.05	ND	NAp	
Fe..	3.00	419	4.562	Responses between the Fe, Pb, As, and Bi peaks are	Fe..	3.00	ND	NAp	
Pb..	.10	284	3.500		K...	NA	65	0.375	
S...	NA	25	.250		Pb..	.10	ND	NAp	
Ti..	.02	49	.562		Si..	>20.0	18	.062	
Zn..	<.10	66	.812		Ti..	.02	ND	NAp	

See notes at end of table.

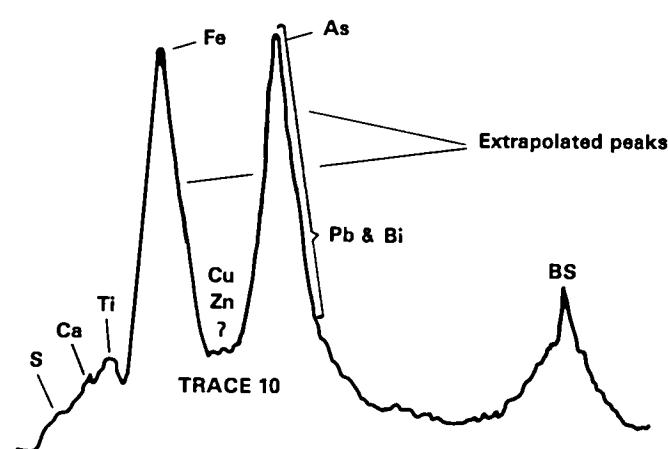
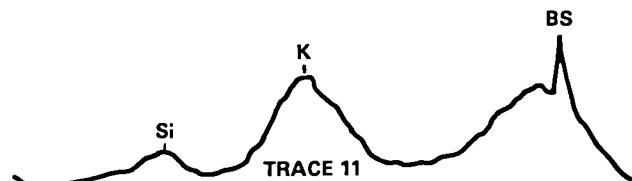
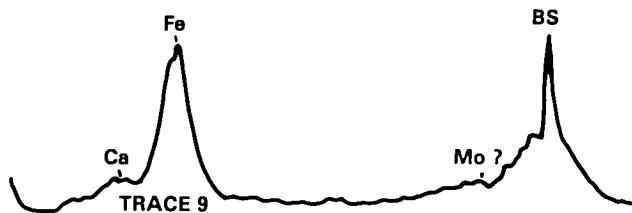


TABLE B-2. - Comparison of minus 200-mesh pulp sample analysis data with PXRFS test results--Continued

Element	Sample analysis, %	PXRFS results		Comments	Element	Sample analysis, %	PXRFS results		Comments
		Total counts	Ratio				Total counts	Ratio	
ANDESITE									
Ca..	10.00	159	1.125	ESA; 180-sec ^{109}Cd irradiation. Fe peak (trace 12) is distorted by the Cu response that shows up as a shoulder to the right of the Fe peak.	Ca..	NA	140	0.550	ESA; 60-sec ^{109}Cd irradiation.
Cu..	2.00	202	1.312		Fe..	NA	440	22.750	
Fe..	7.00	612	6.312		Pb..	0.006	ND	NAp	
Mg..	1.00	ND	NAp		Zn..	32.500	1,512	81.875	
Pb..	.09	ND	NAp		SPHALERITE IN SCHIST				
Rb..	NA	27	.187		MOLYBDENITE IN BULL QUARTZ				
Ti..	.20	ND	NAp		Ca..	NA	35	0.187	ESA; 60-sec ^{109}Cd irradiation.
Zn..	.01	ND	NAp		Fe..	NA	40	.187	
Ca..	10.00	145	2.687	ESA; 60-sec ^{55}Fe irradiation. See trace 13.	Mo..	0.29	103	.562	
Cu..	2.00	ND	NAp		PHOSPHATE ROCK				
Fe..	7.00	ND	NAp		Ca..	26.3000	187	1.500	ESA; 180-sec ^{109}Cd irradiation. Cr, Ti, and V responses are masked by the much stronger Ca response. Much longer analysis time would be required to confirm the presence of Zn and Nb.
Mg..	1.00	ND	NAp		Cr..	.0860	ND	NAp	
Pb..	.09	ND	NAp		Cu..	.0085	21	.187	
Si..	10.00	8	.062		Fe..	.2500	36	.250	
Ti..	.20	ND	NAp		Nb..	NA	26	.187	
Zn..	.01	ND	NAp		S... .	NA	19	.187	
TALC AND CHLORITE, ALTERED DIKE					Sr..	.0990	41	.312	
Ca..	10.57	45	0.437	ESA; 60-sec ^{109}Cd irradiation.	Ti..	.0800	ND	NAp	
Fe..	212.20	297	3.875		V... .	.0290	ND	NAp	
S... .	NA	16	1.870		Zn..	NA	31	.187	

See notes and footnotes at end of table.

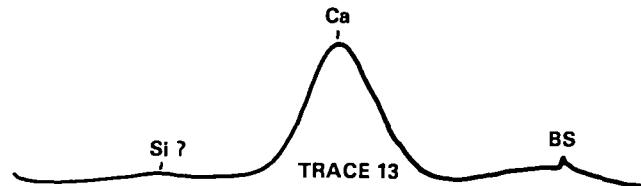
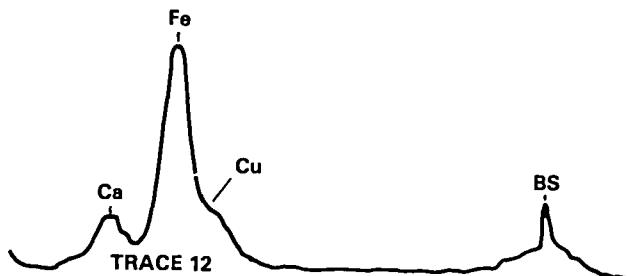


TABLE B-2. - Comparison of minus 200-mesh pulp sample analysis data with PXRFS test results--Continued

Element	Sample analysis, %	PXRFS results		Comments	Element	Sample analysis, %	PXRFS results		Comments
		Total counts	Ratio				Total counts	Ratio	
ALTERED RHYOLITE PORPHYRY									
Ca..	NA	38	0.375	AA; 60-sec ^{109}Cd irradiation.	Ca..	NA	77	0.250	AA; 120-sec ^{109}Cd irradiation.
Cu..	0.0800	ND	NAp		Cu..	0.0010	ND	NAp	
Fe..	NA	104	1.000		Fe..	NA	129	.562	
Mo..	.0006	ND	NAp		Mo..	.0028	ND	NAp	
ANDESITE									
Ca..	NA	47	0.375	AA; 60-sec ^{109}Cd irradiation.	Zn..	.0050	ND	NAp	
Cu..	0.0100	ND	NAp		Ca..	NA	66	.187	AA; 60-sec ^{109}Cd irradiation.
Fe..	NA	121	1.062		Cu..	.0150	ND	NAp	
Mo..	.0035	ND	NAp		Fe..	NA	90	.312	
ARKOSIC SANDSTONE									
Ca..	NA	44	0.500	AA; 60-sec ^{109}Cd irradiation.	Zn..	.0005	ND	NAp	
Cu..	0.0625	ND	NAp		Ca..	NA	73	.187	Do.
Fe..	NA	169	1.275		Cu..	.8010	ND	NAp	
Mo..	.0012	ND	NAp		Fe..	NA	69	.187	
ARGILLITE									
Ca..	NA	49	0.625	AA; 60-sec ^{109}Cd irradiation.	Mo..	.0055	ND	NAp	
Cu..	0.0075	ND	NAp		W...	.0007	ND	NAp	
Fe..	NA	145	1.312		Ca..	NA	58	.250	AA; 90-sec ^{109}Cd irradiation.
Mo..	.0002	ND	NAp		Cu..	.0005	20	.062	
LIMESTONE									
Al..	>4.0	ND	NAp	ESA; 60-sec ^{109}Cd irradiation.	Fe..	NA	50	.187	
Ca..	>10.0	125	2.437		Mo..	<.0002	ND	NAp	
Fe..	3.0	47	.687		W...	<.0002	ND	NAp	
K..	<.06	179	2.687		GRANITE				
Mg..	.6	ND	NAp		Al..	4.000	ND	NAp	ESA; 90-sec ^{109}Cd irradiation.
Ti..	.2	ND	NAp		Au..	NA	20	0.062	
STREAM SEDIMENT FINES					Ca..	.500	69	.250	
Ca..	NA	85	0.750	ESA; 60-sec ^{109}Cd irradiation.	Cb..	.010	ND	NAp	
Cu..	0.0125	ND	NAp		Cu..	.002	ND	NAp	
F...	.0380	ND	NAp		Fe..	1.000	102	.312	
Fe..	NA	1,957	2.687		Ni..	.002	ND	NAp	
Mo..	.0450	ND	NAp		Pb..	.010	ND	NAp	
S...	NA	35	.437		Rb..	NA	65	.125	
W...	.0135	ND	NAp		Ti..	.030	ND	NAp	
Ca..	NA	49	.187	AA; 60-sec ^{109}Cd irradiation.	W...	NA	46	.125	
Cu..	.0005	ND	NAp						
F...	.0260	ND	NAp						
Fe..	NA	34	.125						
Mo..	.0004	ND	NAp						
W...	.0010	ND	NAp						

See notes and footnotes at end of table.

TABLE B-2. - Comparison of minus 200-mesh pulp sample analysis data with PXRFS test results--Continued

Element	Sample analysis, %	PXRFS results		Comments	Element	Sample analysis, %	PXRFS results		Comments
		Total counts	Ratio				Total counts	Ratio	
GRANITE--Continued									
Al..	4.000	ND	NAP	ESA; 60-sec ^{55}Fe irradiation.	As..	0.070	ND	NAP	ESA; 60-sec ^{55}Fe irradiation.
Ca..	.500	ND	NAP		Co..	.020	ND	NAP	
Cb..	3.010	ND	NAP		Cr..	.008	ND	NAP	
Cu..	.002	ND	NAP		Cu..	7.000	ND	NAP	
Fe..	1.000	ND	NAP		Fe..	6.000	ND	NAP	
K...	NA	46	0.375		K...	<.600	12	0.062	
Ni..	.002	ND	NAP		S...	NA	27	.125	
Pb..	.010	ND	NAP		Ti..	.300	ND	NAP	
Si..	>20.000	11	.062		V...	.070	ND	NAP	
Ti..	.030	ND	NAP		Zn..	.030	ND	NAP	
Al..	8.00	ND	NAP	ESA; 180-sec ^{109}Cd irradiation. The Sr response may actually be a Y response--they are adjacent to one another in periodic table and have similar emission energies.	QUARTZ VEIN IN DIORITE--Continued				
Ca..	1.70	119	.375		As..	2.0	349	0.562	ESA; 120-sec ^{109}Cd irradiation.
Cb..	3.02	ND	NAP		Bi..	.2	ND	NAP	
Fe..	4.00	362	1.312		Cu..	.2	ND	NAP	
Pb..	.02	ND	NAP		Fe..	7.0	417	10.187	
Sr..	NA	53	.250		Pb..	>1.0	328	.562	
Y..	.01	ND	NAP		Zn..	.2	ND	NAP	
Al..	8.00	ND	NAP	ESA analysis; 60-sec ^{55}Fe irradiation.	As..	2.0	ND	NAP	ESA; 60-sec ^{55}Fe irradiation.
Ca..	1.70	ND	NAP		Bi..	.2	ND	NAP	
Cb..	3.02	ND	NAP		Cu..	.2	ND	NAP	
Fe..	4.00	ND	NAP		Fe..	7.0	ND	NAP	
K...	NA	75	.562		K...	NA	55	.312	
Pb..	.02	ND	NAP		Pb..	1.0	ND	NAP	
Si..	>20.00	12	.062		Si..	>20.0	10	.062	
Y..	.01	ND	NAP		Zn..	.2	ND	NAP	
QUARTZ VEIN IN DIORITE									
As..	0.070	ND	NAP	ESA; 60-sec ^{109}Cd irradiation.	As..	1.0	883	53.500	AA; 45-sec ^{109}Cd irradiation.
Ca..	NA	85	3.625	As, Co, Cr, Ti, V, and Zn responses all masked by strong Cu and Fe responses.	Ca..	.4	ND	NAP	
Co..	.020	ND	NAP		Cu..	.8	ND	NAP	
Cr..	.008	ND	NAP		Fe..	>10.0	387	25.875	
Cu..	7.000	961	42.000		Pb..	>1.0	ND	NAP	
Fe..	6.000	587	24.812		Zn..	.7	ND	NAP	
Ti..	.300	ND	NAP						
V...	.070	ND	NAP						
Zn..	.030	ND	NAP						

AA Original analysis by atomic absorption. BS Backscatter. ESA Original analysis by mass spectrometer. NA Not analyzed.

NAP Not applicable. ND Not detected or no reading.

¹Analyzed as CaO. ²Analyzed as Fe_2O_3 . ³Analyzed as Cb_2O_5 .

TABLE B-3. - Comparison of USGS sample standards¹ with PXRFS test results

(Ratio value: Total counts divided by backscatter)

Element	Sample analysis, %	PXRFS test results ²		Element	Sample analysis, %	PXRFS test results ²	
		Total counts	Ratio			Total counts	Ratio
Copper...	0.01	21	0.125	Rubidium.	0.010	6	0.062
	.01	25	.125		.021	16	.062
	.05	30	.125		.046	18	.187
	.10	25	.187		.100	31	.250
	.20	54	.312		.215	73	.625
	.50	92	.625		.464	116	1.187
	1.00	130	1.000				
	2.00	301	2.312				
	5.00	945	9.062				
Iron.....	.046	18	.062	Strontium	.001	10	0
	.100	20	.125		.010	13	0
	.220	17	.125		.021	12	.062
	.460	37	.250		.046	13	.125
	1.000	52	.437		.100	16	.250
	2.150	104	1.000		.215	83	.625
	4.640	288	2.375		1.000	267	2.812
	10.000	781	8.250				
Lead.....	.01	10	.062	Zinc.....	.01	27	.125
	.02	14	.062		.02	22	.125
	.05	16	.125		.05	24	.125
	.10	30	.125		.10	26	.187
	.20	28	.187		.20	62	.375
	.50	54	.437		.50	111	.750
	1.00	89	.750		1.00	182	1.437
	2.00	163	1.625		2.00	419	3.125
	5.00	348	4.687		5.00	1,371	14.750
Manganese	.010	376	3.812				
	.022	384	3.875				
	.046	406	4.250				
	.100	404	3.937				
	.215	361	4.187				
	.464	444	4.375				
	1.000	434	5.750				

¹These element standards have a granitic matrix.²120-sec ^{109}Cd irradiation.

TABLE B-4. - PXRFS (^{109}Cd) analysis of selected Bureau of Mines mineral display specimens

Sample and composition (9)	Elements detected by PXRFS	Counts	Element to backscatter ratio	Comments
Carnotite in sandstone; $\text{K}_2(\text{UO}_2)_2(\text{VO}_4)_2 \cdot \text{nH}_2\text{O}$.	K Ca Cu U	77 66 45 213	1.687 1.437 1.375 4.750	30-sec irradiation.
Cinnabar and antimony ¹ ...	Hg S Ca Fe	123 39 51 23	2.312 .250 .812 .437	60-sec irradiation. Sb not detectable with this instrument.
Cupro-tungstate; CuO and WO_3 . ¹	Cu Fe Co Ni Pb Ca	265 170 185 282 535 50	9.375 5.625 6.312 9.187 17.312 1.875	60-sec irradiation. W response is masked by intense Cu response combined with Ni and Co responses.
Erythrite; 37.5% CoO, 38.4% As_2O_5 , 24.1% H_2O .	Co As	189 184	(2) (2)	30-sec irradiation. Overflow indicates a ratio is not possible because backscatter is nonexistent.
Ferberite; 23.7% FeO and 76.3% WO_3 .	Fe W	94 175	(2) (2)	Do.
Galena; 86.6% Pb and 13.4% S.	Pb S Ca Fe	580 42 35 62	56.562 3.500 4.500 5.625	60-sec irradiation.
Garnierite; $(\text{Ni},\text{Mg})\text{SiO}_3 \cdot \text{nH}_2\text{O}$.	K Ni	38 710	3.375 57.000	30-sec irradiation.
Gold telluride ¹	Ca Fe Au	115 24 138	3.437 .875 4.500	60-sec irradiation. Double peak could be from Hg or Pb response as well as Au.
Niccolite; 43.9% Ni and 56.1% As.	Ni As Ca	156 84 60	26.000 12.562 8.000	30-sec irradiation.
Rhodonite; 54.1% MnO and 45.9% SiO_2 .	Mn	671	31.437	45-sec irradiation.
Scheelite in granite; 19.4% CaO and 80.6% WO_3 .	Ca W Fe Rb	93 84 54 4	2.187 1.500 .812 .250	60-sec irradiation.
Scheelite and pyrite; 19.4% CaO, 80.6% WO_3 , 46.6% Fe, 53.4% S.	Ca W Fe	48 78 120	1.062 1.375 2.312	Do.
Uranium in sedimentary rock. ¹	U Ca Fe	106 42 123	.875 .312 1.062	Do.

¹Exact composition percent not known. 20overflow.

APPENDIX C.--NUCLEAR REGULATORY COMMISSION (NRC) LICENSING INFORMATION

All manmade radioactive materials are strictly controlled by the NRC, and it has issued comprehensive regulations for controlling and licensing radioactive isotope sources (called "byproducts" because of their origin in a nuclear reactor).

Briefly, the NRC regulations require the following relative to the PXRFS:

1. The licensing of any company or individual (including Government agencies) that uses isotopes of the strength required in portable X-ray fluorescence spectrometers.
2. The isotopes be under the control of a trained and qualified person at all times.
3. The material be kept reasonably secure against theft or loss by accident.
4. Safety precautions be adequate to protect the user and others in the vicinity from radiation.
5. An authorized individual user be present and directly supervise the use of the spectrometer at any temporary job site.
6. User qualifications include, as a minimum, the completion of the instrument manufacturer's training course.
7. Training by companies is permissible, but the NRC must approve the training course.
8. If multiple users are listed on the license, a radiation protection officer must be named.

"A Guide for Preparation of Byproduct Material Application for the Use of Sealed Sources in Portable and Semiportable Gauging Devices" can be obtained by writing: Materials Licensing Branch, Division of Fuel Cycle and Material Safety, Nuclear Regulatory Commission, Washington, D.C. 20555.





